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The coefficient of first viscosity via three-phonon processes in bulk liquid helium

by

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The coefficient of first viscosity via three-phonon processes in bulk liquid helium

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The contribution of three-phonon processes to the coefficient of first viscosity in bulk liquid helium is evaluated explicitly as a function of temperature, which is shown to have a ${\tt T}^{-1}$ dependence.

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* To whom correspondence should be addressed.



1. Introduction

Since Landau and Khalatnikov's work¹⁾ on transport coefficients, i.e., the thermal conductivity and viscosities of liquid helium, Khalatnikov and Matveev²⁾ reexamined the transport coefficients due to contributions by phonon-roton scattering and noted that the results of Landau and Khalatnikov's work with a normal dispersion deviate from experimental data by as much as a factor of two. The correct anomalous dispersion indicates three-phonon processes (3PP) to be important near absolute zero temperature, and should thus be included in calculations of transport kinetic coefficients.

It is well known that the kinetic equation with temperature gradient for the thermal conductivity vanishes for a pure phonon gas, and thus the corresponding thermal conductivity becomes zero. However, the kinetic equation for the first viscosity does not vanish for a pure phonon gas near zero temperature, and 3PP become important. The purpose of this paper is to evaluate the contribution to the coefficient $\eta_{\rm ph}$ of first viscosity from 3PP which were not included in the work of Landau and Khalatnikov. For small momenta the three-dimensional dispersion relation has been evaluated explicitly by adopting a soft potential to give 3

$$\epsilon(p) = {^{C}_{o}p[1 + \gamma p^{2} - \delta p^{3} + ...]}$$
, (1.1)

where $^{\text{C}}_{\text{O}}$ is the velocity of sound at absolute zero temperature. For bulk liquid helium of density $2.18 \times 10^{-2} \text{ Å}^{-3}$, we can use the following values obtained to fit the experimental data of the excitation energy spectrum:

$$\gamma \mu^2 = 1.51 \text{Å}^2$$
, $\delta \mu^3 = 3.25 \text{Å}^3$, $C_0 = 238.2 \text{ m/s}$, $u = 1.8 \text{ }^6$)

In Sec. 2 we consider phonon-phonon interactions in bulk liquid helium via second quantization to obtain the characteristic time for 3PP. We then solve the kinetic equation for the coefficient of first viscosity in Sec. 3. Finally, in Sec. 4 we discuss numerical results.

2. Phonon-phonon interaction in a three-dimensional quantum liquid

To consider the phonon-phonon interaction in a three-dimensional quantum liquid, we begin with the Hamiltonian H and the internal energy $\mathrm{E}(\rho)^{7}$ for superfluid liquid ⁴He, given by

$$H = \int d^3r \left[\frac{1}{2} \hat{\mathbf{v}} \cdot \hat{\boldsymbol{\rho}} \hat{\mathbf{v}} + E(\boldsymbol{\rho}) \right] , \qquad (2.1)$$

$$E(\rho) = E(\rho_0) + \int d^3p \ n\epsilon(p) , \qquad (2.2)$$

where $\hat{\mathbf{v}}$ and $\hat{\boldsymbol{\rho}}$ are the velocity and density operators expressed in terms of the annihilation operator $\hat{\mathbf{a}}_p$ and creation operator $\hat{\mathbf{a}}_p^{\dagger}$, n is the number density of excitations, and $\hat{\boldsymbol{\rho}}$ is given by $\hat{\boldsymbol{\rho}}_0 + \hat{\boldsymbol{\rho}}'$, the constant and variable parts, respectively. Expanding the internal energy $\mathbf{E}(\boldsymbol{\rho})$ as a function of $\boldsymbol{\rho}'$ and then expressing this expansion in terms of annihilation and creation operators, we obtain

$$H - H_0 + H_3 + H_4 + \dots$$
 (2.3)

$$H_0 = \sum_{p} (\hat{a}_p^{\dagger} \hat{a}_p + \frac{1}{2}) \hbar \omega$$
,

$$\begin{split} & \text{H}_{3} = 4\pi^{3} i (\frac{1}{2V})^{3/2} \sum_{p_{1}} \sum_{p_{2}} \sum_{p_{3}} [(\frac{c_{o}p_{3}}{\rho_{o}p_{1}p_{2}})^{1/2} (\vec{p}_{1} \cdot \vec{p}_{2}) + \frac{\rho_{o}^{2}}{3c_{o}^{2}} \frac{\partial}{\partial \rho} (\frac{c_{o}}{\rho})^{2} \\ & \times (\frac{c_{o}p_{1}p_{2}p_{3}}{\rho_{o}})^{1/2}] [\hat{a}_{p_{1}} \hat{a}_{p_{2}} \hat{a}_{p_{3}} \delta (\vec{p}_{1} + \vec{p}_{2} + \vec{p}_{3}) \exp(-i(\omega_{1} + \omega_{2} + \omega_{3}) t) \\ & + \hat{a}_{p_{1}} \hat{a}_{p_{2}}^{\dagger} \hat{a}_{p_{3}} \delta (\vec{p}_{1} - \vec{p}_{2} - \vec{p}_{3}) \exp(-i(\omega_{1} - \omega_{2} - \omega_{3}) t) + \dots) \\ & + \frac{a_{p_{1}} \hat{a}_{p_{2}}^{\dagger} \hat{a}_{p_{3}} \delta (\vec{p}_{1} - \vec{p}_{2} - \vec{p}_{3}) \exp(-i(\omega_{1} - \omega_{2} - \omega_{3}) t) + \dots) \\ & + \frac{a_{p_{1}} \hat{a}_{p_{2}}^{\dagger} \hat{a}_{p_{3}} \hat{a}_{p_{4}} \delta (\vec{p}_{1} + \vec{p}_{2} + \vec{p}_{3} + \vec{p}_{4}) \exp(-i(\omega_{1} + \omega_{2} + \omega_{3} + \omega_{4}) t) \\ & + \hat{a}_{p_{1}} \hat{a}_{p_{2}}^{\dagger} \hat{a}_{p_{3}} \hat{a}_{p_{4}} \delta (\vec{p}_{1} - \vec{p}_{2} - \vec{p}_{3} + \vec{p}_{4}) \exp(-i(\omega_{1} - \omega_{2} - \omega_{3} + \omega_{4}) t) + \dots] \quad , (2.5) \end{split}$$

where the other six terms to be included in Eq. (2.4) are given by combinations of \hat{a}_{p_1} , \hat{a}_{p_2} , \hat{a}_{p_3} and $\hat{a}_{p_1}^{\dagger}$, $\hat{a}_{p_2}^{\dagger}$, $\hat{a}_{p_3}^{\dagger}$, and fourteen additional terms with the combinations of four creation and four annihilation operators appear in Eq. (2.5).

The nonzero matrix elements of \hat{a}_p and \hat{a}_p^{\dagger} are

$$\langle n_{p}^{-1} | \hat{a}_{p} | n_{p} \rangle = -i (n_{p})^{\frac{1}{2}} e^{i\omega t}$$
,
 $\langle n_{p} | \hat{a}_{p}^{\dagger} | n_{p-1} \rangle = i (n_{p})^{\frac{1}{2}} e^{-i\omega t}$. (2.6)

The matrix element $\langle n_p | H_0 | n_p \rangle$ is $\sum_p (n_p + \frac{1}{2}) M \omega$, where $\omega = c_0 \rho$ and H_0 is thus the Hamiltonian of a simple harmonic oscillator. H_3 and H_4 are the anharmonic perturbative Hamiltonians. Transitions between different states occur due to H_3 , H_4 and other higher-order perturbed Hamiltonians. The transition matrix element from state $|i\rangle$ to state $|f\rangle$ in second-order perturbation theory is given by

$$\langle f|H|i \rangle = \langle f|H_3|i \rangle + \langle f|H_4|i \rangle + \sum_{j} \frac{\langle f|H_3|j \rangle \langle j|H_4|i \rangle}{E_j - E_i}$$
 (2.7)

According to Eqs. (2.6) and (2.4), the first term in Eq. (2.7) contributes only to 3PP, and the second and third terms are related to 4PP.

A finite lifetime of an elementary excitation in a quantum liquid originates from two processes. One is the collision between elementary excitations, and the other is the spontaneous decay of a phonon into two, three or more phonons. Since the collision probability tends to zero at very low temperatures, the collision process will become less important and negligible. At very low temperatures, below 0.6 K, the main excitations in liquid ⁴He are phonons. Therefore, we may consider the decay of the phonon into two phonons only, i.e., 3PP, ⁸⁾ consisting of two processes: the direct process of emission of a phonon, \vec{p}_3 , $\vec{p}_1 = \vec{p}_2 + \vec{p}_3$, and the reverse process of absorption of a phonon, \vec{p}_3 , $\vec{p}_2 + \vec{p}_3 = \vec{p}_1$. The differential decay probability per unit time in three dimensions is given by

$$d\omega = \frac{2\pi}{\aleph} \left| \langle f | H_3 | i \rangle \right|^2 \delta(\epsilon_{f^{-\epsilon}i}) \frac{\nabla^2 d\vec{p}_2 d\vec{p}}{(2\pi \aleph)^6} . \tag{2.8}$$

Since the matrix elements in both processes are given by

$$<\mathbf{f} | \mathbf{H}_{3} | \mathbf{i} >_{\mathbf{D}} - \frac{3!}{2} \frac{(2\pi \mathbf{M})^{3}}{(2\mathbf{V})^{3/2}} \delta(\vec{\mathbf{p}}_{1} - \vec{\mathbf{p}}_{2} - \vec{\mathbf{p}}) \left((\frac{\mathbf{C}_{o}}{\rho_{o}} \frac{\mathbf{p}}{\mathbf{p}_{1} \mathbf{p}_{2}})^{1/2} (\vec{\mathbf{p}}_{1} \cdot \vec{\mathbf{p}}_{2}) \right)$$

$$+ \frac{\rho_{o}^{2}}{3\mathbf{C}_{o}^{2}} \frac{\partial}{\partial \rho} (\frac{\mathbf{C}_{o}^{2}}{\rho}) (\frac{\mathbf{C}_{o}^{\mathbf{p}_{1} \mathbf{p}_{2} \mathbf{p}}}{\rho_{o}})^{1/2} \sqrt{\mathbf{n}_{\mathbf{p}_{1}} (\mathbf{n}_{\mathbf{p}_{2}} + \mathbf{1}) (\mathbf{n}_{\mathbf{p}} + \mathbf{1})} , \qquad (2.9)$$

$$<\mathbf{f}|\mathbf{H}_{3}|\mathbf{i}>_{\mathbf{R}} - \frac{3!}{2} \frac{(2\pi \mathbf{k})^{3}}{(2\mathbf{v})^{3/2}} \delta(\vec{\mathbf{p}}_{1} - \vec{\mathbf{p}}_{2} - \vec{\mathbf{p}}) ((\frac{\mathbf{c}_{o}}{\rho_{o}} \frac{\mathbf{p}}{\mathbf{p}_{1}\mathbf{p}_{2}})^{1/2} (\vec{\mathbf{p}}_{1} \cdot \vec{\mathbf{p}}_{2})$$

$$+\frac{\rho_{o}^{2}}{3C_{o}^{2}}\frac{\partial}{\partial\rho}(\frac{C_{o}^{2}}{\rho})(\frac{C_{o}p_{1}p_{2}p}{\rho_{o}})^{1/2}\}\sqrt{(n_{p_{1}}+1)n_{p_{2}}n_{p}},$$
 (2.10)

where n is the distribution function of phonons with momentum p, the total decay probabilities per unit time of the two processes are

$$\omega_{\rm D} = \frac{\pi c_{\rm o}}{2 \aleph \rho_{\rm o}} ({\rm u} + 1)^2 \int \frac{{\rm d} \vec{p}_2}{(2 \pi \aleph)^3} p_1 p_2 p_1 n_{p_1} (n_{p_2} + 1) (n_p + 1) \delta(\epsilon_{\rm f} - \epsilon_{\rm i}) , \qquad (2.11)$$

$$\omega_{R} = \frac{\pi C_{o}}{2 \mu \rho_{o}} (u+1)^{2} \int \frac{d\vec{p}_{2}}{(2\pi \mu)^{3}} p_{1} p_{2} p(n_{p_{1}}+1) n_{p_{2}} n_{p} \delta(\epsilon_{f} - \epsilon_{i}) , \qquad (2.12)$$

where u is the Grüneisen constant given by $(\rho_{\rm o}/{\rm C_o})(\partial{\rm C_o}/\partial{\rho_{\rm o}})$.

The collision integral, which is the difference between the differential decay probability per unit time for the direct and reverse processes, can be written as

$$J_{3PP} = -\frac{\pi C_{o}}{2 \aleph \rho_{o}} (u+1)^{2} \int \frac{d\vec{p}_{2}}{(2\pi \aleph)^{3}} p_{1} p_{2} p \delta n (n_{p_{1}} - n_{p_{2}}) \delta (\epsilon_{f} - \epsilon_{i}) . \qquad (2.13)$$

where $\delta n = n_p - n_p$, with n_p as the distribution function of phonons with momentum p in the equilibrium state. Since the collision integral $J_{3PP}(n)$ is given by $-(n-n_o)/\tau_{3PP}$, we have

$$\tau_{3PP}^{-1} = \frac{\pi C_{o}}{2 N \rho_{o}} (u+1)^{2} \int \frac{d\vec{p}_{2}}{(2\pi N)^{3}} p_{1} p_{2} p (n_{p_{1}} - n_{p_{2}}) \delta(\epsilon_{f} - \epsilon_{i}) . \qquad (2.14)$$

The three-phonon processes are small-angle processes, and thus p_1 and p_2 are much larger than p_3 . Hence we may use the following relations to evaluate Eq. (2.14):

$$n_{p_1} - n_{p_2} - \left(\frac{\partial n_{p_1}}{\partial \omega_1}\right) \omega , \qquad (2.15)$$

$$\omega_2 - \omega_1 \simeq -\left(\frac{\partial \omega_1}{\partial p_1}\right) p_3 = -\omega \cos\theta$$
 (2.16)

Substitution of Eqs. (2.15)-(2.16) into Eq. (2.14) and integration of Eq. (2.14) over momentum space yields

$$\frac{1}{\tau_{3PP}} = \omega \frac{\pi^3 (u+1)^2 k_B^4 T^4}{30 \mu^3 \rho_0 C_0^5} \left[1 - \frac{60 \pi^2}{7} \gamma (\frac{k_B T}{C_0})^2 \right] . \tag{2.17}$$

3. Viscosity

In this section we shall evaluate the coefficient of first viscosity. We first consider the macroscopic motion of liquid helium, which does not depend on time, with velocity \vec{v} in the z-direction and the gradient directed along x-axis. Then the kinetic equation becomes

$$\vec{\mathbf{v}} \cdot \vec{\nabla} \mathbf{n} = \mathbf{v}_{\mathbf{x}} \frac{\partial \mathbf{n}}{\partial \mathbf{x}} = \mathbf{J}(\mathbf{n}) \quad . \tag{3.1}$$

The phonon distribution function in the motion of liquid helium with velocity \vec{v} is given by

$$n_o = \left[\exp(\frac{Cp - \vec{v} \cdot \vec{p}}{k_B T}) - 1\right]^{-1}$$
, (3.2)

where C is the phonon velocity (equal to C_0 at zero temperature). Substitution of Eq. (3.2) into (3.1) yields

$$n_{o}(n_{o}+1)\frac{Cp}{k_{R}T}\frac{\partial v}{\partial x}\cos\theta\sin\theta\cos\phi = J(n) , \qquad (3.3)$$

where here θ is the angle between \vec{p} and the z-axis. Since 4PP establish energy in the equilibrium state and the distribution function of phonons is related to the variation of phonons, which characterize the chemical potential α , the distribution function becomes

$$n = \left[\exp(\alpha' + \frac{pC}{k_B T} - 1)\right]^{-1} . \tag{3.4}$$

Expanding Eq. (3.4) as a function of α' to obtain the deviation of n from n and considering the linearity of the left-hand side in Eq. (3.3), we can write

$$\delta n = n - n_0 = -n_0(n_0 + 1)\alpha'$$
, (3.5)

$$\alpha' = \alpha \cos \theta \sin \theta \cos \phi$$
 (3.6)

Replacing J(n) by $J_{\mbox{3PP}}(n)$ in Eq. (3.3) and integrating over momentum space, we obtain

$$\int dp \ p^2 J_{3PP}(n) - \cos\theta \sin\theta \cos\phi \left(\frac{k_B T}{c_o}\right)^3 3! \ \zeta(3) \frac{\partial v}{\partial x} . \tag{3.7}$$

Since the collision integral $J_{3PP}(n)$ can be replaced by $-(n-n_o)/\tau_{3PP}$, we can use Eqs. (3.5)-(3.6) together with Eq. (2.17) and $\omega = pC_o/N$ for the evaluation of collision integral:

$$\int dp \ p^2 J_{3PP}(n) = \frac{\pi^3 (u+1)^2 (k_B T)^4}{30 k^4 \rho_o C_o^4} \left(\frac{k_B T}{C_o}\right)^4 3! \zeta(3) \alpha \cos\theta \sin\theta \cos\phi \quad . \quad (3.8)$$

From comparison of Eqs. (3.7) and (3.8) we have

$$\alpha = \frac{30 x^4 \rho_0 c_0^5}{\pi^3 (u+1)^2 (k_B T)^5} \frac{\partial v}{\partial x} - A \frac{\partial v}{\partial x}$$
 (3.9)

The coefficient of first viscosity is obtained from the evaluation of the stress tensor associated with 3PP given by

$$\sigma_{xz} = -\eta_{ph} \frac{\partial v}{\partial x} - \int \frac{d\vec{p}}{(2\pi)!} p_z v_x (n-n_0)$$

$$- \int dp \frac{p^2 \sin\theta d\theta d\phi}{(2\pi)!} p \cos\theta C_0 \sin\theta \cos\phi [-n_0(n_0+1)A \frac{\partial v}{\partial x} \cos\theta \sin\theta \cos\phi] . \qquad (3.10)$$

Performing the integral, Eq. (3.10) becomes

$$\sigma_{xz} = -\frac{3!\zeta(3)}{30\pi^2 k^3} \frac{(k_B T)^4}{c_0^3} A_{\partial x}^{\partial v} . \qquad (3.11)$$

We can get the coefficient of first viscosity by comparing Eq. (3.11) with the first expression in Eq. (3.10):

$$\eta_{\rm ph}(T) = \frac{3! \varsigma(3) k \rho_{\rm o} c_{\rm o}^2}{\pi^5 k_{\rm p} (u+1)^2} \frac{1}{T} = 1.904 \times 10^{-7} \frac{1}{T} .$$
(3.12)

We note that the coefficient of first viscosity $^{1),9)}$ at temperatures below 0.9 K is given by

$$\eta_{\rm ph}(T) = 3.50 \times 10^{-9} \, {\rm T}^{-1/2} \, {\rm e}^{\Delta/k} {\rm B}^{\rm T} (1 + 2.15 \times 10^{-5} {\rm T}^{9/2} {\rm e}^{\Delta/k} {\rm B}^{\rm T})^{-1} , \, {\rm T} < 0.9 \, {\rm K} .$$
(3.13)

4. Numerical Results and discussion

In previous sections we have evaluated the lifetime and the contribution of first viscosity. At relatively low temperatures (< 0.9 K), Eq. (3.13) reduces approximately to

$$\eta_{\rm ph} \sim 1.5 \times 10^{-4} \frac{1}{\rm T}^{5}$$

However, as the temperature approaches absolute zero, the roton density becomes zero, and thus the contribution from the roton-phonon scattering to viscosity will vanish so that the main contribution comes from the 3PP. Therefore, the temperature dependence is changed from T^{-5} to T^{-1} . Figure 1 illustrates the coefficient $\eta_{\rm ph}(T)$ given by Eq. (3.12). The coefficient increases slowly, and near zero temperature it increases very rapidly and finally diverges. In conclusion, we point out that in consideration of the 3PP originating from the anomalous excitation spectrum at low momenta, the temperature variation of first viscosity depends on T^{-1} .

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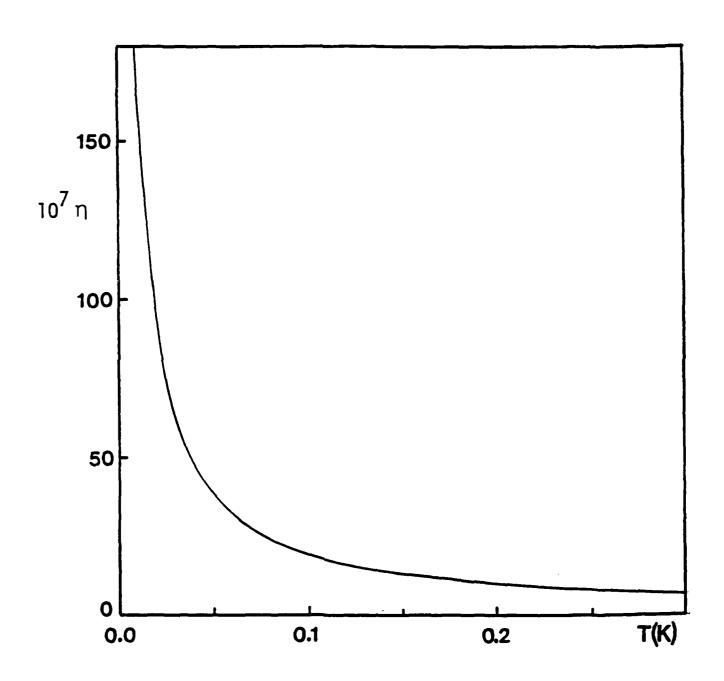
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Figure Caption

Figure 1. Viscosity coefficient due to the contribution from 3PP.



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